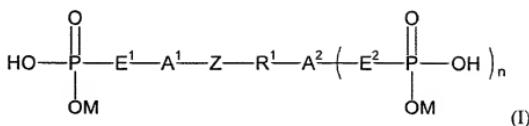


What is claimed is:

1. A compound of formula (I):



where:

A^1 and A^2 are the same or different aryl groups collectively bearing at least one hydrophilic substituent;

E^1 and E^2 are the same or different and are O, S, or NR^2 (where R^2 is a linear or branched $\text{C}_1\text{-C}_{20}$ carbon containing group);

M is H or a pharmaceutically acceptable monovalent cation;

R^1 is a linear or branched, saturated or unsaturated, $\text{C}_1\text{-C}_{20}$ carbon containing group;

Z is a single bond, a carbonyl, CE^3E^4 , or CR^3E^3 , where

E^3 and E^4 are the same or different and are OR^4 , SR^4 , or NR^4_2 , where

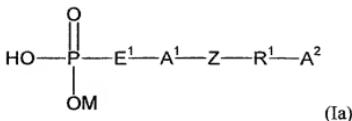
R^3 is a linear or branched $\text{C}_1\text{-C}_{20}$ carbon containing group, and

R^4 is H or a linear or branched $\text{C}_1\text{-C}_{20}$ carbon containing group; and

n is 0 or 1, or a pharmaceutically acceptable salt thereof,

provided that the compound is not 4'-phosphophloretin or a pharmaceutically acceptable salt thereof.

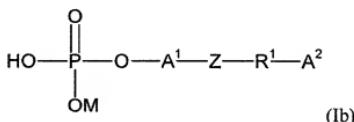
2. The compound of Claim 1 that is a compound of formula (Ia):



where:

A^1 , A^2 , E^1 , M , R^1 and Z are as defined in Claim 1, or a pharmaceutically acceptable salt thereof, provided that the compound is not 4'-phosphophloretin or a pharmaceutically acceptable salt thereof.

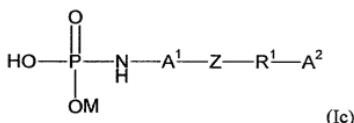
3. The compound of Claim 1 that is a compound of formula (Ib):



where:

A^1 , A^2 , M , R^1 and Z are as defined in Claim 1, or a pharmaceutically acceptable salt thereof, provided that the compound is not 4'-phosphophloretin or a pharmaceutically acceptable salt thereof.

4. The compound of Claim 1 that is a compound of formula (Ic):

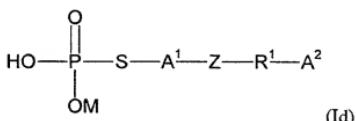


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where:

A¹, A², M, R¹ and Z are as defined in Claim 1, or a pharmaceutically acceptable salt thereof.

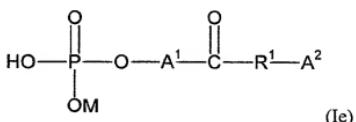
5. The compound of claim 1 that is a compound of formula (Id):



where:

A¹, A², M, R¹ and Z are as defined in Claim 1, or a pharmaceutically acceptable salt thereof.

6. The compound of claim 1 that is a compound of formula (Ie):

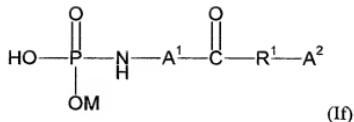


where:

A¹, A², M, and R¹ are as defined in Claim 1, or a pharmaceutically acceptable salt thereof, provided that the compound is not 4'-phosphophloretin or a pharmaceutically acceptable salt thereof.

7. The compound of claim 1 that is a compound of formula (If):

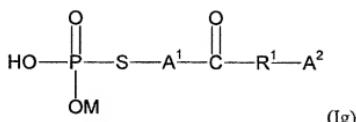
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where:

A^1 , A^2 , M , and R^1 are as defined in Claim 1, or a pharmaceutically acceptable salt thereof.

8. The compound of claim 1 that is a compound of formula (Ig):

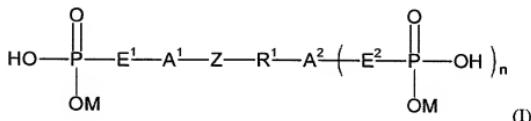


where:

A^1 , A^2 , M , and R^1 are as defined in Claim 1, or a pharmaceutically acceptable salt thereof.

9. The compound of Claim 1 that is 2'-phosphophloretin, 2'-thiophosphophloretin or 2'-aminophosphophloretin or a pharmaceutically acceptable salt thereof.

10. A medication comprising a carrier and a therapeutically effective amount of a compound of formula (I):



where:

A^1 and A^2 are the same or different aryl groups collectively bearing at least one hydrophilic substituent;

E^1 and E^2 are the same or different and are O, S, or NR^2 (where R^2 is a linear or branched C_1-C_{20} carbon containing group);

M is H or a pharmaceutically acceptable monovalent cation;

R^1 is a linear or branched, saturated or unsaturated, C_1-C_{20} carbon containing group;

Z is a single bond, a carbonyl, CE^3E^4 , or CR^3E^3 , where

E^3 and E^4 are the same or different and are OR^4 , SR^4 , or NR^4 , where

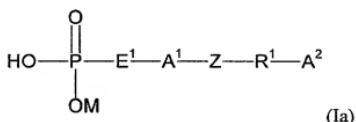
R^3 is a linear or branched C_1-C_{20} carbon containing group, and

R^4 is H or a linear or branched C_1-C_{20} carbon containing group; and

n is 0 or 1,

or a pharmaceutically acceptable salt thereof.

11. The medicament of Claim 10 where the compound is a compound of formula (Ia):

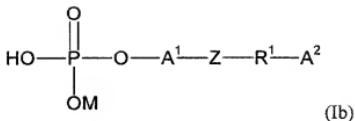


where:

A^1 , A^2 , E^1 , M , R^1 and Z are as defined in Claim 10, or a pharmaceutically acceptable salt thereof.

12. The medicament of Claim 10 where the compound is a compound of formula (Ib):

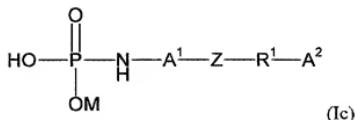
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where:

A^1 , A^2 , M , R^1 and Z are as defined in Claim 10, or a pharmaceutically acceptable salt thereof.

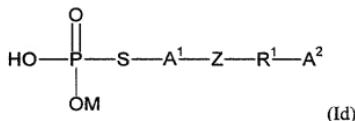
13. The medicament of Claim 10 where the compound is a compound of formula (Ic):



where:

A^1 , A^2 , M , R^1 and Z are as defined in Claim 10, or a pharmaceutically acceptable salt thereof.

14. The medicament of Claim 10 where the compound is a compound of formula (Id):

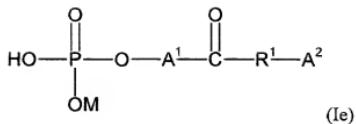


where:

A^1 , A^2 , M , R^1 and Z are as defined in Claim 10, or a pharmaceutically acceptable salt thereof.

15. The medicament of Claim 10 where the compound is a compound of formula (Ie):

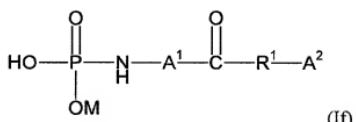
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where:

A^1 , A^2 , M , and R^1 are as defined in Claim 10, or a pharmaceutically acceptable salt thereof.

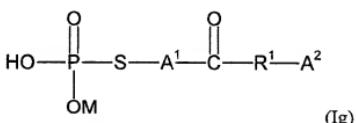
16. The medicament of Claim 10 where the compound is a compound of formula (Ia);



where:

A^1 , A^2 , M, and R¹ are as defined in Claim 10, or a pharmaceutically acceptable salt thereof.

17. The medicament of Claim 10 where the compound is a compound of formula (Ig):

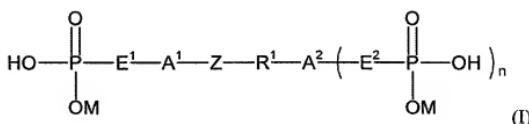


where:

A^1 , A^2 , M , and R' are as defined in Claim 10, or a pharmaceutically acceptable salt thereof.

18. The medicament of Claim 10 where the compound is 2'-phosphophloretin, 2'-thiophosphophloretin or 2'-aminophosphophloretin or a pharmaceutically acceptable salt thereof.

19. A method of inhibiting sodium-mediated phosphate uptake, reducing serum PTH, calcium, calcitriol, or phosphate, or treating renal disease, comprising administration of a compound of formula (I):



where:

A^1 and A^2 are the same or different aryl groups collectively bearing at least one hydrophilic substituent;

E^1 and E^2 are the same or different and are O, S, or NR^2 (where R^2 is a linear or branched C_1-C_{20} carbon containing group);

M is H or a pharmaceutically acceptable monovalent cation;

R^1 is a linear or branched, saturated or unsaturated, C_1-C_{20} carbon containing group;

Z is a single bond, a carbonyl, CE^3E^4 , or CR^3E^3 , where

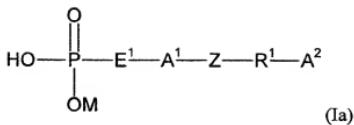
E^3 and E^4 are the same or different and are OR^4 , SR^4 , or NR^4_2 , where

R^2 is a linear or branched C_1-C_{20} carbon containing group, and

R^4 is H or a linear or branched C_1-C_{20} carbon containing group; and

n is 0 or 1, or a pharmaceutically acceptable salt thereof.

20. The method of Claim 19 where the compound is a compound of formula (Ia):



where:

A¹, A², E¹, M, R¹ and Z are as defined in Claim 19, or a pharmaceutically acceptable salt thereof.